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Attorney's Docket No.: 18115-002US1 / SEN-A0123P-US

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

plicant: Minoru Yoshida et al.

Art Unit : 1615

Serial No.: 10/505,380 Examiner: Unknown

Filed

: August 20, 2004

Title

: HISTONE DEACETYLASE INHIBITORS AND METHODS FOR PRODUCING

THE SAME

MAIL STOP AMENDMENT

Commissioner for Patents

P.O. Box 1450

Alexandria, VA 22313-1450

TRANSMITTAL

The following correspondence relating to this application is enclosed for filing:

- 1. Information Disclosure Statement;
- 2. Form PTO-1449;
- 3. Copies of Cited References;
- 4. Copy of the translation of the International Search Report;
- 5. Copy of the translation of the International Preliminary Examination Report; and
- 6. A Return Postcard.

Please date stamp and return the enclosed postcard.

Please apply any charges or credits to Deposit Account No. 06-1050.

Respectfully submitted,

July 12, 200

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INFORMATION DISCLOSURE STATEMENT

Copies of the references listed on the attached form PTO-1449 are enclosed.

This statement is being filed within three months of the filing date of the application or before the receipt of a first Office Action on the merits. Please apply any charges or credits to Deposit Account No. 06-1050.

Respectfully submitted,

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		Applicant Minoru Yoshida et al.	
(Use several st	eets if necessary)	Filing Date August 20, 2004	Group Art Unit 1615

U.S. Patent Documents							
Examiner	Desig.	Document	Publication				Filing Date
Initial	ID	Number	Date	Patentee	Class	Subclass	If Appropriate
	AA 2002/0120099 A1 08/29/2002 Nishino et al.						

	Foreig	n Patent Doo	uments or P	ublished Foreig	n Paten	t Applicat	ions	
Examiner	Desig.	Document	Publication	Country or			Transla	ation
Initial	ID	Number	Date	Patent Office	Class	Subclass	Yes	No
	AB	2 317 003	08/28/2001	Canada				
	AC	1 174 438 A1	01/23/2002	EPO				
	AD	2000256397	09/19/2000	Japan			Abstract	
	AE	2001316283	11/13/2001	Japan			Abstract	*
	AF	2002527449T	08/27/2002	Japan			Abstract	
	AG	2003505417T	02/12/2003	Japan			Abstract	
	AH	WO 00/21979	04/20/2000	WIPO				
	AI	WO 00/52033	09/08/2000	WIPO			Abstract	
	AJ	WO 01/07042	02/01/2001	WIPO				

	Other Documents (include Author, Title, Date, and Place of Publication)				
Examiner	Desig.				
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	AK	Bernhard et al., "Interaction between dexamethasone and butyrate in apoptosis induction: non-additive in thymocytes and synergistic in a T cell-derived leukemia cell line," Cell Death and Differentiation, 1999, 6(7):609-617			
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	AO	Coffey et al., "The Histone Deacetylase Inhibitor, CBHA, Inhibits Growth of Human Neuroblastoma Xenografts in Vivo, Alone and Synergistically with All-Trans Retinoic Acid," Cancer Research, 2001, 61(9):3591-3594			
	AP	Colletti et al., "Broad Spectrum Antiprotozoal Agents that Inhibit Histone Deacetylase: Structure-Activity Relatinships of Apicidin. Part 2," <u>Bioorganic & Medicinal Chemistry Letters</u> , 2001, 11:113-117			
	AQ	Colletti et al, "Design and synthesis of histone deacetylase inhibitors: the development of apicidin transition state analogs," <u>Tetrahedron Letters</u> , 2000, 41:7837-7841			
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Examiner Signature	Date Considered
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Supstitute Form PTO-1449	U.S. Department of Commerce Patent and Trademark Office	Attorney's Docket No. 18115-002US1	Application No. 10/505,380
Information Disclosure Statement by Applicant		Applicant Minoru Yoshida et al.	
(Use several sh	eets if necessary)	Filing Date August 20, 2004	Group Art Unit 1615

		ocuments (include Author, Title, Date, and Place of Publication)
Examiner Initial	Desig. ID	Document
iiiiiai	טו	Dhordain et al., "Corepressor SMRT binds the BTB/POZ repressing domain of the LAZ3/BCL6
	AS	oncoprotein," Proc. Natl. Acad. Sci. USA, 1997, 94:10762-10767
	AT	Dion et al., "Amplification of Recombinant Adenoviral Transgene Products Occurs by Inhibition of Histone Deacetylase," Virology, 1997, 231:201-209
	AU	Ferrara et al., "Histone Deacetylase-targeted Treatment Restores Retinoic Acid Signaling and Differentiation in Acute Myeloid Leukemia," Cancer Research, 2001, 61(1):2-7
	AV	Finnin et al., "Structures of a histone deacetylase homologue bound to the TSA and SAHA inhibitors," Nature, 1999, 401:188-193
	AW	Fischle et al., "A New Family of Human Histone Deacetylases Related to Saccharomyces cerevisiae HDA1p," J. Biol. Chem., 1999, 274(17):11713-11720
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	ABB	He et al., "Distinct interactions of PML-RARα and PLZF-RARα with co-repressors determine differential responses to RA in APL," Nature Genetics, 1998, 18:126-134
	ACC	Hoshikawa et al., "Expression of Differentiation-related Markers in Teratocarcinoma Cells via Histone Hyperacetylation by Trichostatin A," Agric. Biol. Chem., 1991, 55(6):1491-1495
	ADD	Hubbert et al., "HDAC6 is a microtubule-associated deacetylase," Nature, 2002, 417:455-458
	AEE	Inokoshi et al., "Neuronal Differentiation of Neuro 2a Cells by Inhibitors of Cell Cycle Progression Trichostatin A and Butyrolactone I," <u>Biochem. Biophys. Res. Comm.</u> , 1999, 256(2):372-376
	AFF	Ito et al., "p300/CBP-mediated p53 acetylation is commonly induced by p53-activating agents and inhibited by MDM2," EMBO J., 2001, 20(6):1331-1340
	AGG	Juan et al., "Histone Deacetylases Specifically Down-regulate p53-dependent Gene Activation," <u>J. Biol. Chem.</u> , 2000, 275(27):20436-20443
	АНН	Kim et al., "Oxamflatin is a novel antitumor compound that inhibits mammalian histone deacetylase," Oncogene, 1999, 18:2461-2470
	AII	Kim et al., "Histone deacetylases induce angiogenesis by negative regulation of tumor suppressor genes," Nature Medicine, 2001, 7(4):437-443
	AJJ	Komatsu et al., "Cyclic Hydroxamic-acid-containing Peptide 31, a Potent Synthetic Histone Deacetylase Inhibitor with Antitumor Activity," Cancer Research, 2001, 61(11):4459-4466
	AKK	Kwon et al., "Histone Deacetylase Inhibitor FK228 Inhibits Tumor Angiogenesis," <u>Int. J. Cancer</u> , 2002, 97:290-296
	ALL	Li et al., "Causal Relationship between the Loss of RUNX3 Expression and Gastric Cancer," Cell, 2002, 109(1):113-124
	AMM	Lin et al., "Role of the histone deacetylase complex in acute promyelocytic leukaemia," Nature, 1998, 391:811-814
	ANN	Marks et al., "Histone Deacetylase Inhibitors: Inducers of Differentiation or Apoptosis of Transformed Cells," J. Natl. Cancer Inst., 2000, 92:1210-1216

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Substitute Form PTO-1449 (Modified)	U.S. Department of Commerce Patent and Trademark Office	Attorney's Docket No. 18115-002US1	Application No. 10/505,380	
			Applicant Minoru Yoshida et al.	
Ose several sn	eets if necessary)	Filing Date August 20, 2004	Group Art Unit 1615	
(37 CFR §1.98(b)) MADEMARY Other Docum	nents (include Author, 1	Title, Date, and Plac	e of Publication)	

		ocuments (include Author, Title, Date, and Place of Publication)
Examiner Initial	Desig. ID	Document
· -	AOO	Matsuyama et al., "In vivo destabilization of dynamic microtubules by HDAC6-mediated deacetylation," EMBO J., 2002, 21(24):6820-6831
	APP	Meinke et al., "Synthesis of side chain modified apicidin derivatives: potent mechanism-based histone deacetylase inhibitors," Tetrahedron Letters, 2000, 41:7831-7835
	AQQ	McKinsey et al., "Signal-dependent nuclear export of a histone deacetylase regulates muscle differentiation," Nature, 2000, 408:106-111
	ARR	Minucci et al., "A histone deacetylase inhibitor potentiates retinoid receptor action in embryonal carcinoma cells," Proc. Natl. Acad. Sci. USA, 1997, 94(21):11295-11300
	ASS	Munster et al., "The Histone Deacetylase Inhibitor Suberoylanilide Hydroxamic Acid Induces Differentiation of Human Breast Cancer Cells," Cancer Research, 2001, 61(23):8492-8497
	ATT	Nakajima et al., "FR901228, a Potent Antitumor Antibiotic, Is a Novel Histone Deacetylase Inhibitor," Exp. Cell Res., 1998, 241(1):126-133
	AUU	Nan et al., "Transcriptional repression by the methyl-CpG-binding protein MeCP2 involves a histone deacetylase complex," Nature, 1998, 393(6683):386-389
	AVV	Petti et al., "Complete remission through blast cell differentiation in <i>PLZF/RARa</i> -positive acute promyelocytic leukemia: in vitro and in vivo studies," <u>Blood</u> , 2002, 100(3):1065-1067
	AWW	Primeau et al., "Synergistic Antineoplastic Action of DNA Methylation Inhibitor 5-AZA-2'-Deoxycytidine and Histone Deacetylase Inhibitor Depsipeptide on Human Breast Carcinoma Cells, Int. J. Cancer, 2003, 103:177-184
AX	AXX	Saito et al., "A synthetic inhibitor of histone deacetylase, MS-27-275, with marked in vivo antitumo activity against human tumors," Proc. Natl. Acad. Sci. USA, 1999, 96(8):4592-4597
	AYY	Verdel and Khochbin, "Identification of a New Family of Higher Eukaryotic Histone Deacetylases, J. Biol. Chem., 1999, 274(4):2440-2445
	AZZ	Verdel et al., "Active maintenance of mHDA2/mHDAC6 histone-deacetylase in the cytoplasm," Current Biology, 2000, 10:1-3
	AAAA	Wang et al., "Inhibitors of Histone Deacetylase Relieve ETO-mediated Repression and Induce Differentiation of AML1-ETO Leukemia Cells," Cancer Research, 1999, 59(12):2766-2769
	ABBB	Yang et al., "Isolation and Characterization of cDNAs Corresponding to an Additional Member of the Human Histone Deacetylase Gene Family," J. Biol. Chem., 1997, 272(44):28001-28007
	ACCC	Yoshida et al., "Potent and Specific Inhibition of Mammalian Histone Deacetylase Both in Vivo an in Vitro by Trichostatin A," <u>J. Biol. Chem.</u> , 1990, 265(28):17174-17179
	ADDD	Yoshida et al., "Trichostatin A and trapoxin: novel chemical probes for the role of histone acetylation in chromatin structure and function," <u>BioEssays</u> , 1995, 17(5):423-430
	AEEE	Yoshida et al, "Effects of Trichostatins on Differentiation of Murine Erythroleukemia Cells," <u>Canc</u> <u>Research</u> , 1987, 47(14):3688-3691

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